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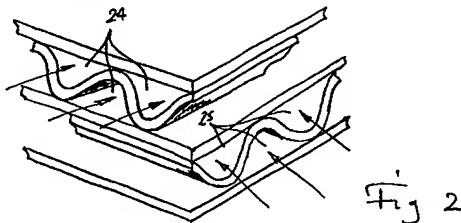
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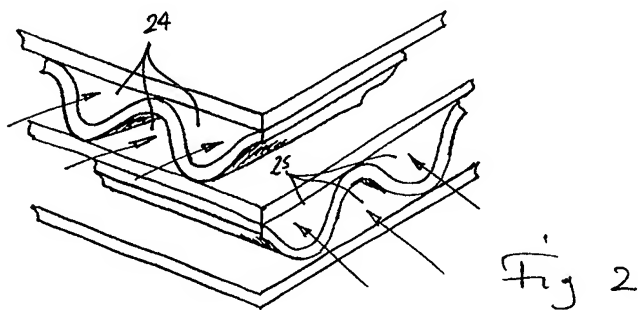
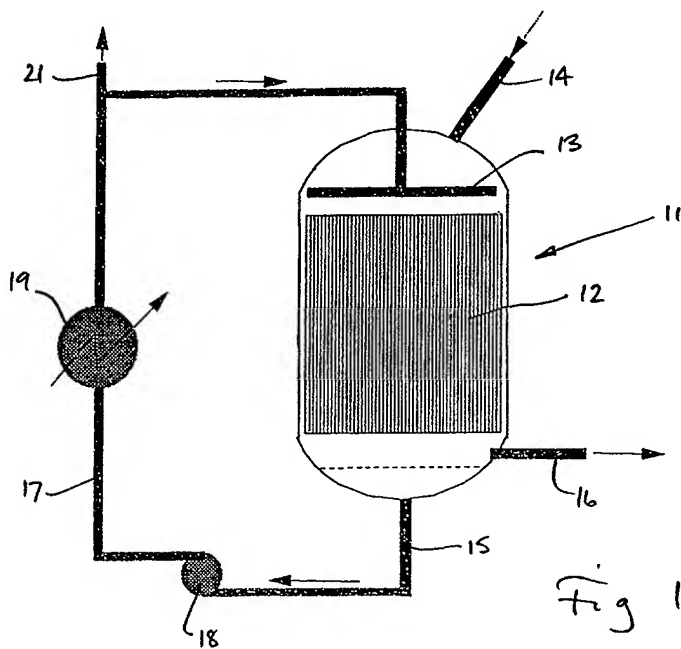
(54) Abstract Title  
Fischer-Tropsch reactor

(57) The reactor comprises a monolithic solid body arranged in discrete and continuous channels. The walls of the channels 24 contain a conventional Fischer-Tropsch catalyst; conventional promoters may also be included. The body preferably also comprises channels for conducting coolants 25 which are arranged transversely to the reaction channels.

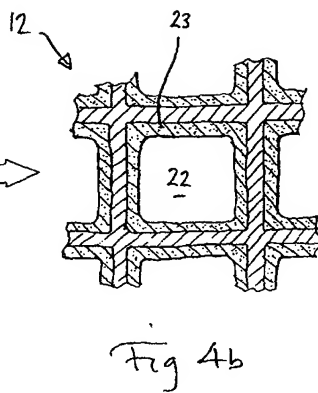
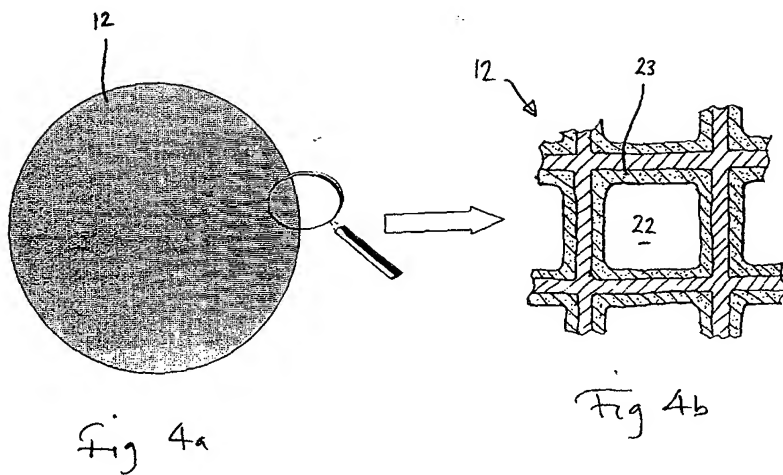
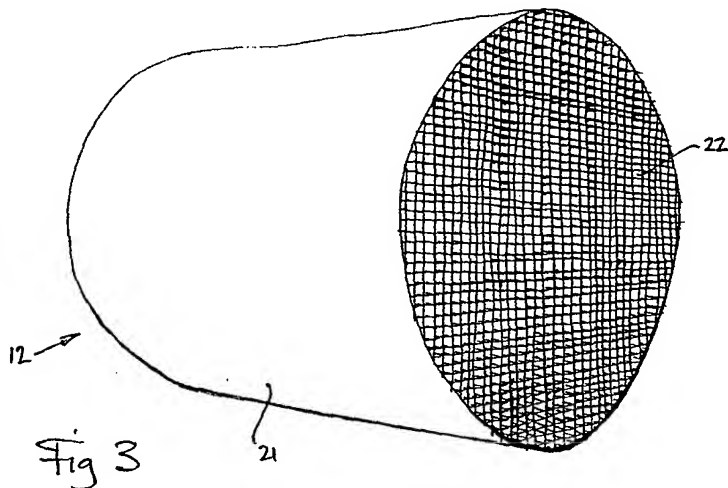


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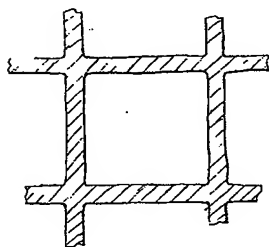


Fig 5a

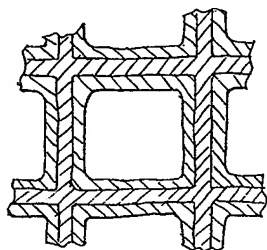
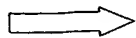


Fig 5b

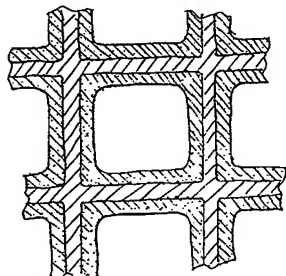


Fig 5c

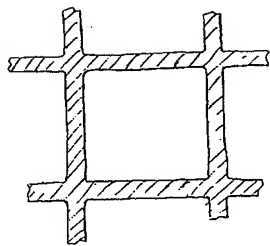


Fig 6a

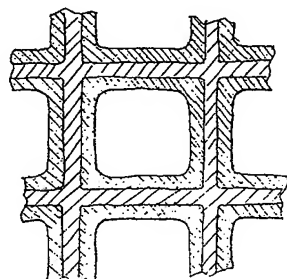
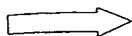


Fig 6b

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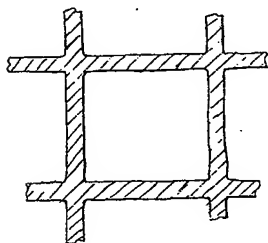


Fig 7a

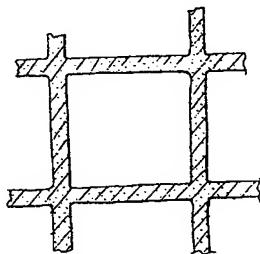
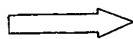


Fig 7b

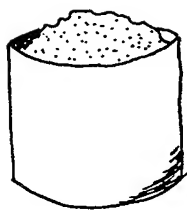


fig 8a

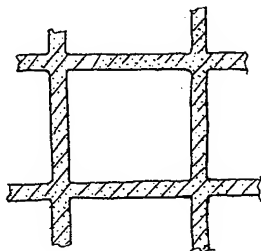
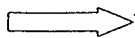
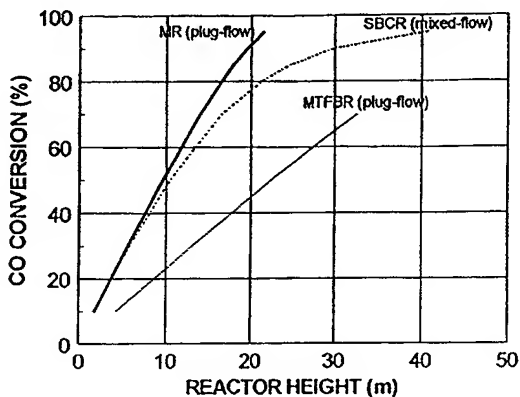


Fig 8b

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Fig 9

Comparison of reactors.  $P = 25$  bar,  $H_2/CO = 2$ .



SBCR = Slurry Bubble Column Reactor

(Solid fraction = 0.15 (vol),  $u_g = 16$  cm/s)

MR = Monolith Reactor

(Solid fraction = 0.15 (vol),  $u_g = 16$  cm/s,  $k = k_{SBCR}$ )

MTFBR = Multitubular Fixed-Bed Reactor

(Solid fraction = 0.6 (vol),  $u_g = 50$  cm/s,  $k = k_{SBCR}/3$ ).

Fischer-Tropsch Synthesis

The present invention relates to a Fischer-Tropsch synthesis.

5       Conventional methods of operating a low temperature Fischer-Tropsch synthesis employ fixed bed reactors. The catalytically active material is generally carried on relatively large carrier particles and this results in poor intraparticle mass transfer characteristics. Also,  
10       the catalyst particles are packed in the tubes of a shell and tube arrangement with coolant on the shell side. This means that the space/time yield is limited by the heat transfer in the catalyst bed.

15       An alternative is to operate the fixed bed as a trickle-bed reactor. This may have advantages from the point of view of heat transfer, however, the poor intraparticle mass transfer characteristics remain.

20       It has also been proposed to carry out an F-T synthesis in a slurry bubble column reactor. Here, the catalyst particles would be significantly smaller since they would be in suspension in the liquid product. This in turn gives rise to significantly improved mass transfer characteristics within the catalyst particles. However, this type of reaction system can be  
25       disadvantageous in that separation of the catalyst from the product can be troublesome. There is also back mixing, due to the nature of the reactor, which is less efficient in terms of reactor volume than a plug flow reactor.

30       It is an object of the present invention to provide a reaction system for an F-T synthesis which has high mass transfer characteristics at the catalyst and in which heat transfer is not a significantly limiting factor, without the drawback of a difficult catalyst

separation.

As world oil resources diminish, natural gas is becoming more attractive as an energy source and methods of upgrading this to higher hydrocarbon fuels are increasing in importance. Thus, in one system, the methane (natural gas) is reformed to CO and H<sub>2</sub> and this synthesis gas is subjected to a Fischer-Tropsch reaction to form higher hydrocarbon products.

Gas fields are now being discovered at significant distances offshore and under certain circumstances it is not commercially viable to pipe the gas on-shore for processing. It would be advantageous, therefore, if the natural gas in these remote fields could be processed on board marine vessels at the fields and then taken to port by those vessels or by other transport vessels. Under these conditions, reactor size is crucial, and the existing fixed bed and slurry reactors are undesirably large. They are also sensitive to movement, particularly the slurry reactor which would be unstable in heavy seas.

It is therefore a further object of the invention to provide a reaction system for an F-T synthesis in which the reactor size is minimised and which is less sensitive to movement than existing systems.

Accordingly, the invention is directed to the use of a monolithic catalyst to conduct a Fischer-Tropsch synthesis, in which the monolith comprises a solid body defining a series of discrete and continuous channels extending from one end of the body to the other, the walls of the channels consisting of or containing a Fischer-Tropsch catalyst.

The invention also provides a method of conducting a Fischer-Tropsch synthesis reaction which comprises: passing synthesis gas comprising H<sub>2</sub> and CO through discrete and continuous channels in a monolithic



catalyst, the walls of the channels consisting of or containing a Fischer-Tropsch catalyst; removing the liquid product from the monolith; and removing heat produced in the reaction in the liquid product.

5            Preferably heat from the reaction is removed from the liquid product stream outside the reactor and a portion of the liquid product stream is recycled to the reactor. Unreacted synthesis gas may be recycled from the reactor, for example to the synthesis gas production  
10 unit.

            Preferably, the synthesis gas feed and the liquid product flow co-currently. Preferably the synthesis gas feed and liquid product travel along the channels in a slug flow or Taylor Flow regime. Taylor Flow of a gas  
15 and liquid in a channel is defined as periodic cylindrical gas bubbles in the liquid having almost the same diameter as the channel and without entrained gas bubbles between successive cylindrical bubbles. Preferably, the flow is downwards.

20           Alternatively, the gas/liquid flows could be counter-current.

            In a preferred form, the invention provides a reaction system for a Fischer-Tropsch synthesis which comprises a reactor including a monolithic catalyst and  
25 having an inlet for synthesis gas comprising  $H_2$  and CO and an outlet for liquid product, the monolithic catalyst comprising a solid body defining a series of discrete and continuous channels extending from one end of the body to the other, the walls of the channels consisting of or  
30 containing a Fischer-Tropsch catalyst, whereby the synthesis gas is supplied via the inlet and is passed through the channels where the synthesis takes place and liquid product is removed via the outlet, the heat produced by the reaction being removed from the system by

the liquid product.

Mass transfer (diffusion) effects are also very important in determining selectivity. Diffusion rates in the liquid phase are typically 3 orders of magnitude  
5 slower than in the gas phase, meaning that even slow reactions may be diffusion limited in liquid phase. Recent investigations have shown that a moderate diffusion resistance within the catalyst pellet (intraparticle) can give a strong negative influence on  
10 C5+ selectivity. The key parameter is the characteristic diffusion distance, determined by the catalyst pellet size or the thickness of a catalytic layer. As a guideline, negative effects on selectivity are experienced for diffusion lengths above 0.1-0.4 mm, corresponding to 0.2-0.8 mm diameter spherical pellets  
15 (the exact value depending on catalyst properties and reaction conditions).

Conventional fixed bed reactors typically use 2-6 mm diameter pellets in order to avoid an unacceptable pressure drop through the catalyst bed. In a fixed bed  
20 reactor, the selectivity problem can be solved by using catalyst pellets where the catalytic material is deposited in a thin outer layer ("egg-shell" catalysts). However, this means that only a fraction of the catalyst present in the reactor is participating in the reaction,  
25 reducing the solid fraction from typically 60% to about 23% for a 2 mm diameter spherical particle with a 0.15 mm catalytic layer. In a slurry reactor, the selectivity problem is solved by using small catalyst particles, usually 0.1 mm or less.  
30

In the proposed monolith reactor concept, a short diffusion distance (typically  $<0.15$  mm) can be maintained without having to reduce the fraction of active material, as a result of the fact that the catalyst is located in

the thin walls of the monolith structure.

5       The Fischer-Tropsch synthesis is a strongly  
exothermal reaction and effective heat transfer is a  
prerequisite for successful reactor operation. In a  
15       fixed bed reactor, the catalyst is located within tubes  
and the heat is removed by steam generation on the shell  
side. The space time yields are therefore limited by the  
heat transfer properties of the reactor. In addition,  
the maximum conversion per pass is limited by the high  
10       gas velocities necessary for achieving optimum heat  
transfer. The heat transfer properties can be improved  
by decreasing the tube diameter, but this increases the  
pressure drop. In addition, the cost and weight of the  
reactor will increase strongly with decreasing tube  
15       diameter. In practice, significant axial and radial  
temperature gradients are unavoidable in fixed bed  
reactors used for FT-synthesis.

      In slurry reactors, the catalyst/slurry is located  
on the shell side and the heat of reaction is removed by  
20       steam generation on the tube side. Due to the turbulent  
motion of the slurry, the heat transfer properties are  
more favourable and the necessary heat transfer area is  
drastically reduced when compared to a fixed bed reactor.

      In the proposed monolith reactor design, cooling is  
25       performed by direct heat removal by the production stream  
(preferably the heavy FT products) which may circulate.  
The circulating liquid can then be cooled in an external  
heat exchanger. If necessary, the cooling can be carried  
out in stages by dividing the reactor in different  
30       sections with separate cooling circuits.

      In slurry reactors, the productivity of the reactor  
(space time yield) may be limited by the gas-liquid mass  
transfer, i.e. from gas bubbles to the liquid. The  
characteristic mass-transfer area is therefore the gas-

liquid interfacial area, which is roughly an order of magnitude smaller than the external catalyst area.

5 In a monolithic reactor operating in two-phase flow (gas + liquid), particularly under Taylor Flow conditions, mass transfer occurs mainly in the thin film between the cylindrical bubbles and the channel walls containing the catalytic material. This mode of flow occurs over a wide range of gas and liquid superficial velocities. However, at very high gas velocities, the  
10 flow regime changes to "annular flow", characterised by less favourable mass transfer properties.

The high gas-liquid mass transfer rates (compared to trickle-bed reactors or slurry reactors) and the high liquid velocities possible (compared to trickle-bed  
15 reactors) are factors that make the monolith type of reactor well suited for FT-synthesis.

Another desirable property of monolith reactors is the extremely low pressure drop for liquids and gases over the catalyst bed, which is 1-2 orders of magnitude  
20 lower than in a packed bed (trickle-bed or gas-phase fixed-bed) reactor. This is an important factor at high gas and liquid superficial velocities and in cases where reactants or liquid are recycled to the reactor. The problem of uneven gas or liquid distribution, bypassing  
25 etc. is also much smaller than in packed-bed reactors.

A primary goal in any FT-process is to achieve a high conversion per pass, in order to achieve high yields and reduce recycle and investment costs. For any reaction, the reaction kinetics and the mixing  
30 characteristics of the reactor will determine the necessary reaction volume for reaching the desired conversion. For a given feedrate to the reactor, an increase in conversion is achieved by increasing the height of the reactor.

A slurry reactor is characterised by extensive backmixing, resulting in relatively flat concentration profiles for reactants and products. As an approximation, the reactor can be assumed to operate as a perfectly mixed reactor.

On the other hand, tubular fixed-bed reactors and a monolith reactor will show negligible backmixing, i.e. near plug-flow. The narrow channels of a monolith and the Taylor-flow mode of operation in two-phase flow results in almost perfect plug-flow. Scale-up is therefore simple since the entire reactor can be described by a single channel.

Classical chemical reaction engineering principles teach that chemical reactions characterised by a positive order dependence of reactant concentrations are most efficiently carried out in plug-flow. In other words, the necessary reaction (or catalyst) volume (i.e. the reactor height) is smaller for a reactor operating in plug-flow, compared to mixed flow. The disadvantage of mixed-flow reactors increase with increasing conversion and increasing reaction order.

If the FT-synthesis is assumed to have a 1. order dependency on hydrogen partial pressure, the necessary volume for a perfectly mixed reactor will be approximately 3 times larger than in a plug-flow reactor at 90% conversion and the same catalyst concentration, temperature and pressure. However, 1. order kinetics is not an accurate representation of the FT-reaction. The apparent pressure order is less than 1.0, and the kinetics can for the present purpose be approximated by a power function:

$$r = k P_{H_2}^{0.5} P_{CO}^{-0.1}$$

$r$  = CO consumption rate

$k$  = rate constant

$P_{H_2}$ ,  $P_{CO}$  = partial pressure of  $H_2$  and CO, respectively

5 A comparison of the performance of various FT-reactors is shown in Figure 9. The monolith and slurry reactors have been assumed to have the same catalyst concentration per volume of reactor and a typical inlet gas velocity  $u_g = 16$  cm/s. The fixed-bed reactor, due to the less favourable  
10 heat transfer properties, has been assumed to operate at 1/3 of the rates achievable in the SBCR or MR cases, either as a result of lower catalyst activity, lower temperatures or both. It is well known from FT and methanol synthesis that multi-tube fixed bed reactors  
15 must operate at high gas velocities in order to obtain optimum heat transfer. 50 cm/s has been chosen as a representative value for the fixed bed reactor. The reaction rates and inlet velocities are representative values for the reactors considered.

20 It is evident from Figure 9 that the difference in reactor height for the SBCR and MR is relatively small for low-moderate conversions (up to about 60%). However, for conversions above 80% and in particular above 90%, the difference in reactor height becomes significant.  
25 For the fixed bed reactor, the high linear velocity makes it virtually impossible to achieve high single pass conversions.

Thus, the monolith can achieve higher conversions than a backmixed slurry reactor, and is of particular  
30 interest for cases where extremely high conversions are desired (>90%).

As mentioned above, a slurry reactor needs continuous catalyst/product separation in order to operate successfully and for producing a catalyst-free

product. Such arrangements are not necessary in a monolith reactor. Heavy products are easily removed from the liquid recycle stream as required for maintaining a constant inventory of liquid.

5 In many monolith applications, the thermal stability of the material and the ability to withstand rapid temperature variations are both of great importance. Therefore, the channel structure of a monolith usually consists of a low-surface area ceramic material. The  
10 surface area can be increased by depositing a high-surface area material (like  $\gamma\text{-Al}_2\text{O}_3$ ), e.g. by the so-called washcoating technique. Catalytically active materials can then be incorporated into the washcoat by known techniques, like impregnation, precipitation, ion-exchange, vapour deposition etc. Alternatively, the low  
15 surface area base material can be washcoated with the catalytic material itself.

Thus in one preferred form, the monolithic catalyst comprises an inactive substrate with a relatively low  
20 specific surface area, and, lining the channels, a relatively high specific area catalyst support impregnated with a catalytically active material. Preferably the catalyst support material and the active material are deposited simultaneously on the walls of the  
25 channels. Alternatively, the catalyst support material is first deposited on the walls of the channels and is subsequently impregnated with the active material. The inactive substrate may be a ceramic material or a metal. Examples of suitable materials are set out in Table 1  
30 (taken from "Monolithic Catalysts for Nonautomobile Applications" by S Irandoust and B Andersson, Catal. Rev. Sci. Eng., 30(3), 1988).

Table 1  
Materials for Monolithic Substrates

Name	Composition
$\alpha$ - and $\gamma$ -Alumina	$Al_2O_3$
Cordierite	$2MgO \cdot 2Al_2O_3 \cdot 5SiO_2$
Cordierite-mullite	$2MgO \cdot 2Al_2O_3 \cdot 5SiO_2 - 2Al_2O_3 \cdot 2SiO_2$
Magnesium aluminate-spinel	$MgO - MgO \cdot Al_2O_3$
Mullite	$3Al_2O_3 \cdot SiO_2$
Mullite-aluminum titanate	$3Al_2O_3 \cdot 2SiO_2 - Al_2O_3 \cdot TiO_2$
Silica	$SiO_2$
Silicon carbide	$SiC$
Silicon nitride	$Si_3N_4$
Spinel	$MgO \cdot Al_2O_3$
Titania	$TiO_2$
Zeolites	$Al_2O_3 - SiO_2$
Zirconia	$ZrO_2$
Zirconia-spinel	$ZrO_2 - MgO \cdot Al_2O_3$
Metallic	$Fe - Cr - Al - Yt$

A monolith-based FT catalyst can thus be made by impregnation (or by other techniques) of an active FT-metal (Co, Fe, Ru, Ni) and suitable promoters on a high-surface area washcoated monolith or by washcoating the finished FT-catalyst onto the low-surface area monolith.

In slower reactions like the Fischer-Tropsch synthesis, thermal stability is not a critical factor. In such cases, monoliths can be made directly from high surface area materials, for example gamma- $Al_2O_3$ ,  $SiO_2$ ,  $TiO_2$  or zeolites. The catalytic material (e.g. cobalt, iron ruthenium or nickel in the case of FT-synthesis) and optionally suitable promoters can then be incorporated into the total volume of the monolith (by any of the known techniques), thus increasing the catalyst loading of the reactor compared to the washcoat method.



Alternatively, the monolith can be produced directly from the FT-catalyst. Production of high surface area monoliths is usually achieved by extrusion.

Thus, in another preferred form, the monolithic catalyst comprises an inactive substrate with a relatively high specific surface area in which the walls of the channels are impregnated with a catalytically active material. In an alternative form, the monolithic catalyst comprises an extrusion of a relatively high specific surface area material incorporating a catalytically active material. Preferably, the high specific surface area material is  $\gamma$ - $\text{Al}_2\text{O}_3$ ,  $\text{SiO}_2$ ,  $\text{TiO}_2$  or zeolite. Generally suitable materials also include transition aluminas, alumina/silica, magnesium aluminate spinel and titania/silica, or any high surface area ceramic material.

The particular F-T active catalyst material selected will depend to an extent on the final product required, however, preferred materials include Co, Fe, Ru and/or Ni. The catalyst may include a promoter. Suitable promoters include Re, Pt, Ir, Rh, Pd and Ru.

The monolith preferably has an open area or void fraction of 50 to 90%, more preferably 60-80%, for example 70. It may have a cell density of 100 to 1000 cells/in<sup>2</sup> (15.5 to 155 cells/cm<sup>2</sup>), preferably 200 to 600 cells/in<sup>2</sup> (31.0 to 93.0 cells/cm<sup>2</sup>), most preferably 300-500. The wall thickness is preferably 0.05 to 0.4 mm, more preferably 0.1-0.3, for example 0.15.

The monolith, which may be extruded, may have a length in the centimetres to metres range, the ultimate maximum length being limited only by manufacturing techniques. A reactor of the required size can be produced by stacking together monolithic catalyst blocks.

The invention also extends to a monolithic catalyst

per se, as described, for carrying out an F-T synthesis.

The monolithic catalyst reactor system described exhibits in particular the following qualities, many of which represent significant advantages over known F-T reactor systems:-

- 5 a) Plug-flow, very high conversions possible or short reactors for moderate - high conversions
- b) Simple scale-up
- c) Short diffusion distance in monolith walls, good
- 10 selectivity
- d) High gas-liquid mass-transfer rates in two-phase flow
- e) High liquid and gas throughputs possible
- f) Low pressure drop
- 15 g) Temperature control by direct cooling of catalyst with liquid medium and external heat removal
- h) No wax-catalyst separation necessary

20 The invention may be carried into practice in various ways and some embodiments will now be described by way of example with reference to Figures 1 to 8 of the accompanying drawings, in which:-

Figure 1 is a schematic diagram of a reaction system according to the invention;

25 Figure 2 is a perspective view in part of an alternative arrangement;

Figure 3 is an isometric sketch of a monolithic catalyst;

Figure 4a is an end view of the monolithic catalyst;

30 Figure 4b is an enlarged view of part of Figure 4a;

Figures 5a to 5c are sequential views, similar to Figure 4b, in the manufacture of a monolithic catalyst according to a first method;

Figures 6a and 6b are sequential views showing a

second method;

Figures 7a and 7b are sequential views showing a third method;

5        Figures 8a and 8b are sequential views showing a fourth method; and

Figure 9 is a comparison in graph form of the performance of various FT-reactors.

10        Figure 1 shows a reaction system for an F-T synthesis embodying the present invention. The system comprises a reactor 11 containing a monolithic catalyst 12 and a liquid distributor 13 above the catalyst monolith 12. The reactor 11 also has a gas inlet 14 near the top, a liquid outlet 15 at the base and a gas outlet 16 near the base. The liquid outlet 15 is joined to the  
15        liquid distributor via a recycle stream 17 which includes a pump 18 and a heat exchanger 19. The recycle stream 17 has a liquid product outlet 21.

20        In use, synthesis gas ( $H_2$  and CO) is introduced via the gas inlet 14 and product liquid recycled to the distributor 13. The liquid and gas are drawn down through longitudinal channels in the monolith 12 under a Taylor flow regime. The gases react to form hydrocarbons, with a preference for hydrocarbons with 5 or more carbon atoms. The liquid product is withdrawn  
25        via the outlet 15 by means of the pump 18 and is cooled in the heat exchanger 19 where water is heated and vaporised. A portion is removed at the product outlet 21 and the remainder is recycled to the distributor 13. Any unreacted gas and any gaseous product is removed at the  
30        gas outlet 16 and possibly recycled.

In an alternative construction shown in Figure 2, the monolith is a cross-flow design. In this case, the gases are arranged to travel along one series of channels 24 and the cooling liquid along the transverse channels

25 which alternate vertically with the gas channels.

The monolith 12 as shown in Figures 3, 4a and 4b is cylindrical, though any suitable shape could be employed such as rectangular. It comprises an outer wall 21, preferably formed of the same material as the monolith body which is formed with regular longitudinal channels 22 of square section though, again, any shape could be employed. The channels 22 are discrete and continuous and extend from one end of the monolith to the other. In this case the monolith 12 is of low surface area cordierite, the surface area being 0.1 to 1.0 m<sup>2</sup>/g. The walls of the channels 22 have a layer 23 of a high surface area material, such as  $\gamma$ -Al<sub>2</sub>O<sub>3</sub>, the surface area being about 200 m<sup>2</sup>/g. This material is impregnated with an active catalyst material such as Co with a promoter such as Re.

One method of manufacturing the monolith 12 is shown in Figures 5a to 5c. A monolithic support is formed from an inactive low surface area (typically 0.1 to 1.0 m<sup>2</sup>/g) material such as cordierite by mixing fine cordierite powder with plasticisers and binders to form a paste which is then extruded. The extruded body is dried and fired at a high temperature and has a large number of square section channels (Figure 5a).

Active material is then applied to this monolith support by a washcoating technique. A high surface area (20 to 200 m<sup>2</sup>/g) material such as  $\gamma$ -alumina may be used. The monolith is dipped into a slurry of the high surface area material and this is then dried and heat treated, leaving a layer of the high surface area material on the walls of the channels (Figure 5b).

Active material such as Co is introduced into the washcoat by impregnation, precipitation ion exchange from solution or other known methods (Figure 5c).

An alternative washcoat method is shown in Figures 6a and 6b. In this case the monolith support is prepared in the same way as that shown in Figures 5a (Figure 6a). Then, however the monolith is dipped in a liquid suspension containing a fine powder of a Fischer-Tropsch catalyst. Thus when the washcoated layer is dried, it already contains the active material (Figure 6b).

Another method of producing the monolith is shown in Figures 7a and 7b. Here, the support material is itself an inactive high surface area material. The monolithic support is therefore formed by extrusion in the same way as described above, but using the high surface area material (Figure 7a).

The support material is then impregnated with the active material from solution in the same way as the high surface area layer in the washcoating methods (Figure 7b).

In an alternative direct extrusion method, shown in Figures 8a and 8b, a paste of catalyst (i.e. a high surface area support material already containing active material) is extruded, dried and fired to form a high surface area support already containing the active material.

Naturally, to a certain extent, these methods could be combined in order to produce the desired catalyst properties.

The invention will now be further illustrated in the following Examples.

#### Catalyst preparation

##### Example 1

##### *Conventional catalysts*

465 g of  $\gamma\text{-Al}_2\text{O}_3$  (Puralox B-5207 from Condea) was sieved

to 38-53  $\mu\text{m}$  particles and calcined in air at 500°C for 10 hours before use. 582 g of  $\text{Co}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$  and 9.7 g of  $\text{HReO}_4$  (80-85% solution) was dissolved in deionized water to produce 740 ml of impregnating solution which was mixed with the  $\gamma\text{-Al}_2\text{O}_3$  support. The resulting catalyst was dried in air at 95°C for 2 days before calcination in air at 300°C for 16 hours. The calcined catalyst contained 17 wt% Co and 1 wt% Re. A portion of the catalyst was pressed to larger pellets and then crushed and sieved to various particle sizes, namely 53-75  $\mu\text{m}$  (designated catalyst "A"), 75-150  $\mu\text{m}$  (catalyst "B") and 425-850  $\mu\text{m}$  (catalyst "C").

#### Example 2

##### *Washcoated monolith catalysts*

Another portion of the catalyst described above was wet grinded further in a ball mill and sieved to <38 microns for use in preparing monolithic catalysts. Approximately 10 cm long cylindrical low surface area cordierite monoliths (Celcor (reg. trademark) from Corning) with a diameter of about 0.9 cm were used as base materials. The monoliths had a cell density of 400 cells/in<sup>2</sup> (62 cells/cm<sup>2</sup>) and 76% open area (void fraction). A slurry of the  $\text{CoRe}/\text{Al}_2\text{O}_3$  catalyst was prepared by mixing 50 g of the <38 micron catalyst with 100 ml of water. The cordierite monolith was then washcoated with the  $\text{CoRe}/\text{Al}_2\text{O}_3$  catalyst by dipping the monoliths into the catalyst slurry and blowing away excess slurry from the monolith channels gently with an air gun. 2 catalyst samples were produced in this manner, designated "D" and "E" after dipping the monolith 1 and 2 times respectively. In the latter case, the excess slurry was removed and the catalyst was dried between the first and second dip. The monoliths were dried in air at 130°C for 16 hours before calcination in

air at 400°C for 2 hours. The calcined monolith catalysts "D" and "E" contained 0.3 g and 0.7 g of CoRe/Al<sub>2</sub>O<sub>3</sub> washcoat, respectively.

5      **Example 3**

*High surface area monolith catalysts*

10       $\gamma$ -Al<sub>2</sub>O<sub>3</sub> monoliths from Corning (based on Vista Catapal (reg. trademark) alumina from Vista Chemical Company) with a cell density of 400 cells/in<sup>2</sup> (62 cells/cm<sup>2</sup>) and 70% open area was used as a base material. The surface area of the monolith was 145 m<sup>2</sup>/g and the pore volume was 0.53 cm<sup>3</sup>/g. Two 10 cm long cylindrical monoliths with 1 cm diameter were used for catalyst preparation. Two identical catalyst samples were produced by dipping the

15       $\gamma$ -Al<sub>2</sub>O<sub>3</sub> monoliths into a solution containing 100 g of Co(NO<sub>3</sub>)<sub>2</sub>·6H<sub>2</sub>O, 1.75 g of HReO<sub>4</sub> (80-85% solution) and 50 g of deionized water. After removing the excess solution, the monoliths were dried in air at 95°C for 16 hours before calcination in air at 400°C for 12 hours. The calcined

20      catalysts contained 12 wt% Co and 0.6 wt% Re. The first sample (designated "F") was used without modification, while the second sample ("G") was crushed and sieved to 38-150  $\mu$ m particles.

25      Catalyst testing

The catalysts were tested in a 20 cm long 1 cm ID stainless steel fixed-bed reactor closely surrounded by an electrically heated aluminium jacket. The conventional catalysts (powder) were diluted with an inert material

30      (non porous SiC) in a 1:5 weight ratio in order to minimise temperature gradients. The monolithic catalysts were used as prepared. All catalysts were reduced in flowing hydrogen at a space velocity of 5000 cm<sup>3</sup> (STP)/(g<sub>cat</sub>.h) at atmospheric pressure at 350°C for 16

hours (heating rate from ambient temperature: 1 K/min). The space velocity refers to the amount of CoRe/Al<sub>2</sub>O<sub>3</sub> catalyst present in the reactor. After reduction, the catalyst was cooled to 170°C in flowing H<sub>2</sub> and purged with He before increasing the pressure to 13 bar (1.3 MPa) and switching to a feed mixture consisting of synthesis gas (H<sub>2</sub>/CO = 2.1) premixed with 3 mol% N<sub>2</sub> as internal standard. The reaction temperature was then slowly increased to the desired temperature (usually 210°C). On-line samples were taken at regular intervals and analysed for N<sub>2</sub>, CO, CO<sub>2</sub> and C<sub>1</sub>+ hydrocarbons on a gas chromatograph.

#### Catalyst performance

The performance of the catalysts is shown in the Table 2 below. All data are averaged over a period of >10 hours and are representative of stabilised catalysts, i.e. after more than 30 hours on stream.

Catalysts D-F are according to the invention. Catalysts A-C and G are only included for comparison purposes.

It is shown that the washcoated monolith catalysts (D&E) are as active as the conventional (powder) catalysts (A-C). The application of two washcoat layers (cat. E) on the cordierite base does not influence the catalyst performance compared to the single layer monolith (cat. D).

The high-surface area monolith catalyst (F) contained more cobalt per volume of reactor than the other catalysts and it was found that 195°C was the maximum acceptable reactor temperature due to heat removal limitations in a gas-phase reactor. These limitations



will not be present in a monolith reactor operating with liquid coolant. After correcting for the temperature difference, it is evident that the high-surface area monolith is at least as active per unit mass of cobalt as the conventional (powder) catalysts and also similar to the crushed high-surface area monolith catalyst (G). The high C5+ selectivity indicates that hot-spots were not present during reaction.

Although the small particle conventional catalysts (A&B) show catalyst performances (reaction rates and selectivities) on the same level as the monolith based catalysts, it is known that such particle sizes cannot be used in a commercial fixed-reactor due to an unacceptable pressure drop. Particle sizes  $> 1 \text{ mm}$  ( $> 1000 \mu\text{m}$ ) are necessary in commercial fixed-bed reactors. However, the results for the largest particles in the present series of catalysts (425-850  $\mu\text{m}$ , cat. C) show that conventional catalysts of this size produce unacceptably high methane and light gas ( $\text{C}_2\text{-C}_4$ ) selectivities. The  $\text{CO}_2$  selectivity is also higher than for small particle or monolith-based catalysts. These differences are caused by intraparticle diffusion effects. Under the prevailing conditions, the catalyst pores become filled with liquid hydrocarbon products and large concentration gradients will be present for the reactants ( $\text{H}_2, \text{CO}$ ) in catalyst particles of sufficient size. Due to the difference in the transport properties of  $\text{H}_2$  and  $\text{CO}$ , the particle  $\text{H}_2/\text{CO}$ -ratio will increase towards the particle centre and cause the loss of C5+ selectivity. Due to the particular reaction kinetics of  $\text{CO}$  hydrogenation on cobalt catalysts (i.e. the  $\text{H}_2$  and  $\text{CO}$  partial pressure dependency of the reaction rate), strong diffusion effects on selectivity are observed before any significant effects are observed on

the overall hydrocarbon formation rate. This is in agreement with the data in the Table 2.

Table 2

- 5 Catalyst test results. P = 13 bar (1.3 MPa),  $H_2/CO = 2.1$ , 3% inerts in feed. The space velocities and reaction rates are based on the weight of active catalyst (CoRe/ $Al_2O_3$ ).

CATALYST	Temp. (°C)	GHSV (h <sup>-1</sup> ) <sup>7</sup>	CO conv. (%)	Reaction rate		Selectivity (%)			
				( $g_{HC}/g_{cat} \cdot h$ )	( $g_{HC}/g_{Co} \cdot h$ )	CH <sub>4</sub>	C <sub>2</sub> -C <sub>4</sub>	C <sub>3</sub> +	CO <sub>2</sub>
A (53-75 micron powder)	210	10.900	27.7	0.61	3.6	9.0	7.4	82.9	0.7
B (75-150 micron powder)	210	10.100	26.2	0.54	3.2	9.1	7.5	82.7	0.8
C (425-850 micron powder)	210	7.300	33.0	0.49	2.9	21.5	12.7	64.4	1.5
D (Cordierite base, 0.3 g cat.)	210	16.800	16.3	0.56	3.3	8.2	7.3	83.8	0.7
E (Cordierite base, 0.7 g cat.)	210	8.400	29.1	0.50	2.9	8.9	6.5	84.0	0.6
F (High surf.area monolith)	195	4.060	27	0.22 (0.5) <sup>**</sup>	1.8 (4.0) <sup>**</sup>	7.1	6.2	86.4	0.3
G (Crushed high surf.area monolith)	210	6.000	30	0.36	2.9	8.2	7.1	84.5	0.2

<sup>7</sup> cm<sup>3</sup> (STP) syngas/g<sub>cat</sub>·h

<sup>\*\*</sup> Estimated reaction rate at 210°C using an activation energy of 100 kJ/mol

## Claims:

1. The use of a monolithic catalyst to conduct a Fischer-Tropsch synthesis, in which the monolith comprises a solid body defining a series of discrete and continuous channels extending from one end of the body to the other, the walls of the channels consisting of or containing a Fischer-Tropsch catalyst.
2. A method of conducting a Fischer-Tropsch synthesis reaction which comprises: passing synthesis gas comprising  $H_2$  and CO through discrete and continuous channels in a monolithic catalyst, the walls of the channels consisting of or containing a Fischer-Tropsch catalyst; removing the liquid product from the monolith; and removing heat produced in the reaction.
3. A method as claimed in Claim 2, in which heat produced in the reaction is removed in the liquid product.
4. A method as claimed in Claim 3, in which heat is removed from the liquid product stream outside the reactor and a portion of the liquid product stream is recycled to the reactor.
5. A method as claimed in any of Claims 2 to 4, in which heat produced in the reaction is removed by cross-current flow of a cooling medium in separate channels through the body of the monolithic catalyst.
6. A method as claimed in Claim 4, in which the synthesis gas feed and the liquid product flow co-currently.

7. A method as claimed in Claim 6, in which the synthesis gas feed and liquid product travel along the channels in a Taylor Flow regime.
- 5 8. A method as claimed in Claim 7, in which the synthesis gas feed and liquid product flow downwards through the monolith.
- 10 9. A method as claimed in Claim 4, in which the synthesis gas feed and the liquid product flow counter-currently.
- 15 10. A reaction system for a Fischer-Tropsch synthesis which comprises a reactor including a monolithic catalyst and having an inlet for synthesis gas comprising  $H_2$  and CO and an outlet for liquid product, the monolithic catalyst comprising a solid body defining a series of discrete and continuous channels extending from one end of the body to  
20 the other, the walls of the channels consisting of or containing a Fischer-Tropsch catalyst, whereby the synthesis gas is supplied via the inlet and is passed through the channels where the synthesis takes place and liquid product is removed via the outlet, and heat  
25 produced by the reaction is removed from the system by the liquid product.
- 30 11. A system as claimed in Claim 10, in which the monolithic catalyst comprises an inactive substrate with a relatively low specific surface area, and, lining the channels, a relatively high specific area catalyst support containing a catalytically active material.
12. A system as claimed in Claim 11, in which the

catalyst support material and the active material are deposited simultaneously on the walls of the channels.

5 13. A system as claimed in Claim 11, in which the catalyst support material is first deposited on the walls of the channels and the active material is subsequently incorporated into the support material.

10 14. A system as claimed in any of Claims 11 to 13, in which the substrate is a ceramic material or a metal.

15 15. A system as claimed in Claim 14, in which the ceramic material is alpha-alumina or cordierite.

16 16. A system as claimed in Claim 10, in which the monolithic catalyst comprises an inactive substrate with a relatively high specific surface area in which the walls of the channels contain a catalytically active material.

20 17. A system as claimed in Claim 10, in which the monolithic catalyst comprises an extrusion of a relatively high specific surface area material incorporating a catalytically active material.

25 18. A system as claimed in any of Claims 11 to 17, in which the relatively high specific surface area material is  $\text{Al}_2\text{O}_3$ ,  $\text{SiO}_2$ ,  $\text{TiO}_2$  or zeolite.

30 19. A system as claimed in any of Claims 10 to 18, in which the active Fischer-Tropsch catalyst material is Co, Fe, Ru and/or Ni.

20. A system as claimed in any of Claims 10 to 19,

further incorporating a promoter.

21. A system as claimed in Claim 20, in which the promoter is Re, Pt, Ir, Rh, Pd and/or Ru.

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22. A system as claimed in any of Claims 10 to 21, characterised in that the monolithic catalyst has a void fraction of 50 to 90%.

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23. A system as claimed in any of Claims 10 to 22, characterised in that the monolithic catalyst has a cell density of 100 to 1000 cells/in<sup>2</sup> (15.5 to 155 cells/cm<sup>2</sup>).

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24. A system as claimed in any of Claims 10 to 23, characterised in that the monolithic catalyst has a wall thickness of 0.05 to 0.4 mm.

25. A system as claimed in any of Claims 10 to 24, comprising a plurality of the monolithic catalysts.

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**Patents Act 1977**  
**Search Report under Section 17**

**Databases searched:**

UK Patent Office collections, including GB, EP, WO & US patent specifications, in:

UK Cl (Ed.O): B1E; C5E (ECE, ECF)

Int Cl (Ed.6): B01J 35/04

Other: ONLINE: WPI, CLAIMS

**Documents considered to be relevant:**

Category	Identity of document and relevant passage	Relevant to claims
A	GB 1385907 (ICI) page 1 lines 38-63; page 3 lines 97-111	
A	EP 0298943 A2 (SVENSKA EMISSION. AB) column 3 lines 38-41; column 4 lines 28-38	

X Document indicating lack of novelty or inventive step  
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